

Identification and Physical Mapping of A Polymorphic Human T Cell Receptor V β Gene with a Frequent Null Allele

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Summary

Germline variation in genes that encode the human T cell receptors (TCRs) may have an important influence in shaping the immune T cell repertoire. In this report we describe a frequent null allele of the human V β 18 gene, resulting from a nucleotide substitution that creates a stop codon (CGA \leftrightarrow TGA). Approximately 11% of the population tested was homozygous for this null allele, indicating that this is a frequent "hole in the repertoire." We confirmed that there is a greatly reduced (undetectable) level of V β 18 mRNA in peripheral blood lymphocytes from an individual homozygous for this null allele. In addition, all heterozygous individuals expressed detectable levels of only the functional V β 18 allele in their peripheral blood lymphocytes. Two other DNA polymorphisms were identified in V β 18, one of which would result in an amino acid substitution in an expressed V β 18 gene. Genotypes for all three of these V β 18 DNA polymorphisms were determined in a group of unrelated individuals. Statistical analyses of the associations between alleles of the V β 18 polymorphisms and those of other DNA polymorphisms in the TCR β locus suggested a close physical proximity between the V β 18 gene and the 3' end of the C β 2 region. This localization of human V β 18 had been previously predicted by the sequence homology between human V β 18 and mouse V β 14, a V gene segment previously mapped to 3' of the mouse C β genes. We confirmed this localization of the human V β 18 gene by isolating a cosmid clone that contains both the V β 18 and C β 2 gene segments. Mapping by restriction enzyme digestion and by the polymerase chain reaction indicated that the V β 18 gene segment is approximately 9 kb 3' of the C β 2 gene, making this the only known human V β gene 3' of the C β region.

A diverse repertoire of TCRs is generated by a variety of somatic mechanisms, including the random rearrangement of gene segments, N region diversification, and junctional diversity. In addition to these somatic effects, allelic variations arising from DNA polymorphisms in the TCR genes also contribute to germline diversity in the population (1). In the past, RFLPs detected with probes from TCR V genes were used as indirect measures of DNA polymorphism within the V genes (2–6). More recently, however, direct sequence analysis has demonstrated the presence of germline DNA polymorphisms in the TCR that result in amino acid substitutions. Specifically, Robinson (7) has reported a polymorphism in the V β 1 gene segment that results in a change from glutamine to histidine, while Li et al. (8) reported two polymorphisms in the V β 6.7 gene that result in a serine to arginine and also a glycine to glutamic acid substitution. Although the ultimate impact of these or other allelic variations on the TCR repertoire remains to be determined, the

analysis of TCR polymorphisms may provide a greater understanding of the potential relationship between DNA polymorphisms in the TCR and susceptibilities to certain autoimmune diseases (reviewed in references 1 and 9).

To further assess the extent of TCR germline polymorphism, we have compared the sequence of TCR V gene segments from multiple individuals. In this study, we report a striking example of a DNA polymorphism in the V β 18 gene that results in the introduction of a termination codon (the V β 18 gene has also been referred to as the V β 20 gene in another nomenclature by Kimura et al. [9a]). 11% of the individuals analyzed were homozygous for this null allele, and therefore could not express a functional V β 18 protein product. Analysis of V β 18 expression using RNA from PBL confirmed this relationship between V β 18 RNA message and the stop codon. Additionally, we have also obtained the entire genomic sequence of the V β 18 gene, and have determined its physical location to be at the 3' end of the C region genes.

This localization is opposite of other known human V β genes, but is syntenic with the location of a homologous TCR V gene segment (V β 14) previously mapped in the mouse.

Materials and Methods

Human DNA and RNA. A panel of DNAs isolated from the lymphoblastoid cell lines of six unrelated Caucasians was used for polymorphism screening, and was kindly provided by R. A. Gatti. The DNA of 76 Caucasian parents and certain families from Centre d'Etude du Polymorphisme Humaine (CEPH) (10), pedigrees were used for polymorphism typing. For the V β 18 expression analysis, RNA was isolated from PHA-stimulated PBL from several small families (kindly provided by R. Spielman, University of Pennsylvania, Philadelphia, PA) using the acid guanidinium thiocyanate-phenol-chloroform extraction method (11).

Oligonucleotides. Amplification primers and ligation probes were synthesized using a DNA synthesizer (384 or 391; Applied Biosystems, Inc., Foster City, CA). The sequences of amplification primers and ligation probes used in polymorphism typing of the V β 18 gene segment are shown in Table 1. The 5' allele-specific ligation probes were biotinylated as previously described (12), and the adjoining 3' reporter probes were phosphorylated using "5'-phosphate-on" according to the manufacturer's instructions (Clontech, Palo Alto, CA). Ligation probes were purified by reverse-phase HPLC, and the phosphorylated 3' reporter probes were enzymatically labeled with digoxigenin as previously described (13).

DNA Amplification. DNA samples were amplified by the PCR. For determining the V β 18 genotypes by restriction enzyme digestion, PCR was performed using a DNA thermal cycler (Perkin-Elmer Cetus, Emeryville, CA) in a reaction mix containing reaction buffer (Amersham Corp., Arlington Heights, IL), nucleotides (final concentration, 20 μ M each), DNA polymerase (Hot Tub; Amersham Corp.) (final concentration, 10 U/ml), 0.3 μ M each of primer V β 18-3 and V β 18-4 (Table 1), and 10–100 ng of genomic DNA. The amplification profile was 35–40 cycles of 1 min at 94°C, 1 min at 60°C, and 2 min at 72°C.

For determining the V β 18 genotypes by OLA, DNA samples were amplified in a 96-well microtiter plate thermal cycler (MJ Research, Watertown, MA). Genomic DNA (10 ng) was mixed with a standard PCR buffer (10 mM Tris-HCl, pH 8.3, 50 mM KCl, 1.5 mM MgCl₂, and 0.001% gelatin) containing 40 μ M of each of the four deoxynucleotide triphosphates, 0.5 μ M of each primer, and 10 U/ml of Taq polymerase (Perkin-Elmer Cetus) (14). The amplification profile was 40 cycles of 20 s at 93°C, 40 s at 55°C, and 90 s at 72°C.

For PCR-based mapping of the H29 cosmid, 50 ng of cosmid DNA was mixed with the same mixture of PCR buffer, nucleotides, and enzyme as described directly above for amplifying DNA in a 96-well format. The primer pair used for amplification was 5'-CTGTTTCCCTGAAGATTGAG-3' (3' end of C β 2) and 5'-ACTGGGTGGAGGATATTTGTAAG-3' (3' end of V β 18). This mix was then subjected to 40 cycles of 1 min at 93°C, 1 min at 55°C, and 2 min at 72°C in a DNA thermal cycler (Perkin-Elmer Cetus). The amount of time at 72°C was extended by 5 s after each cycle and is essential to amplifying larger DNA products.

DNA Sequencing. For purposes of DNA sequencing, a large-scale PCR (200 μ l) was performed according to the conditions described above. To eliminate an additional manipulation, mineral oil was not used during the amplification so that ethanol precipitation was done in the same tube as used for amplification. The concentrated PCR products were purified by electrophoresis through 1% low-melting point agarose (Bethesda Research Laboratories,

Gaithersburg, MD), and the amplified products excised from the gel. 10 μ l of the PCR product in the gel slice was used directly as the template for sequencing in 96-well assay plates using the Sequenase sequencing kit (U.S. Biochemical Corp., Cleveland, OH) (15). The sequencing primers used were the same as those used for the original amplification. Sequence from the V β 18 gene region was obtained using the H29 cosmid. Sequence-specific oligonucleotide primers were used to sequence across both strands of the cosmid template (\sim 2 μ g of cosmid DNA template per labeling reaction). Sequencing was done using the Sequenase sequencing kit, and the annealing and radiolabeling steps were performed at 37°C (15).

Restriction Digestion Analysis. Restriction digestion of the PCR products was performed without further purification of the DNA. 10 μ l of the amplified DNA was digested with 10 U of either KpnI or Sau3A (Bethesda Research Laboratories). Only the addition of BSA (final concentration, 100 μ g/ml) was necessary for complete digestion with KpnI, whereas addition of the manufacturer's 10 \times digestion buffer was required for digestion with Sau3A. Restriction enzyme-digested PCR products were subjected to electrophoresis in a 3% agarose gel (including 2% Nusieve agarose; FMC Bioproducts, Rockland, ME) containing ethidium bromide, and visualized by UV illumination.

Oligonucleotide Ligation Assay (OLA).¹ Analyses of amplified DNA samples using primers V β 18-1 and -2 for each of the V β 18 polymorphisms were performed using an OLA detection system previously described in detail (13, 14). In this assay, each sample was analyzed by a ligation reaction for each allele. Each reaction contained one of the 5' biotinylated probes specific for that allelic form of the polymorphism, the common 3' reporter probe labeled with digoxigenin, an aliquot of the amplified DNA target, and T4 DNA ligase. If the probes were hybridized to a perfectly matched target, the biotinylated allele specific and reporter probes were joined by the ligase. To determine whether the ligation reaction had been successful, the reaction was transferred to a streptavidin plate to capture the biotinylated probe. The plate was washed and then assayed for the presence of the covalently attached reporter molecule using an enzyme-linked immunoassay system (13).

Cosmid Identification and Restriction Digest Mapping. A human cosmid library constructed in a modified pWE15 vector (16) and containing 15 genome equivalents was screened for V β 18. The cosmid H29 was identified by hybridization (final stringency was 0.5 \times SSC at 65°C) to a PCR-amplified V β 18 gene segment that had been cloned into the M13 phage. The H29 cosmid was mapped with EcoRI, HindIII, BamHI, and XbaI using a standard protocol of partial digestion, and then sequential hybridization with vector-specific oligonucleotides (final stringency wash of 6 \times SSC at 65°C), and to TCR C β and V β 18 probes.

V β 18 RNA Expression. For PCR analysis of RNA, cDNA was synthesized from 1–2 μ g of total RNA by random hexamer priming using a cDNA synthesis kit (Amersham Corp.). Approximately 25 ng of the cDNA was PCR amplified with 25 cycles using a 5' V β 18 and 3' C β together with a 5' and 3' C α primer pair, as listed in Fig. 6, b and c, legends. The products were subjected to electrophoresis at 150 V through 2% agarose and vacuum blotted with 0.4 N NaOH onto Genetran nylon membrane (Plasco, Marlboro, MA). The membrane was hybridized overnight to a C β and C α DNA probe that were ³²P radiolabeled separately by random priming. These probes had been combined as a 50:50 mixture based on the amount of incorporated ³²P. The hybridized blot was then

¹ Abbreviation used in this paper: OLA, oligonucleotide ligation assay.

washed several times up to a final stringency of $0.1 \times$ SSC at 65°C . The PCR products were detected by autoradiography.

Results

leader region, through the first intron and the V β 18 gene segment, and ends within the 3' noncoding sequence. The 3' noncoding sequence also includes the recombination signal (heptamer-nonamer sequences) for somatic DNA rearrangement. One of the interesting features of the human V β 18 gene sequence is its similarity to mouse V β 14 (19). This similarity across species is 77% in the coding regions, and 63% overall.

Genotyping of the *Vβ18* Polymorphisms. Genotypes for the

Figure 1. Complete genomic DNA and amino acid sequence of the human $V\beta 18$ gene segment. The DNA and amino acid sequence polymorphisms are shown at positions 524, 620, and 661. The actual DNA sequence of the three $V\beta 18$ haplotypes observed is indicated by the A, B, and C lines (-, identical sequence). The heptamer-nonamer somatic recombination signals found in the 3' noncoding region are italicized and bolded. PCR primers used to amplify this gene segment are underlined. The double-underlined DNA sequences were found to be different compared with the original reported cDNA sequence of HUT102 (20), and were not found to represent additional polymorphism. These sequence data are available from EMBL/GenBank/DBJ under accession number Z33967 HSTCRVB18.

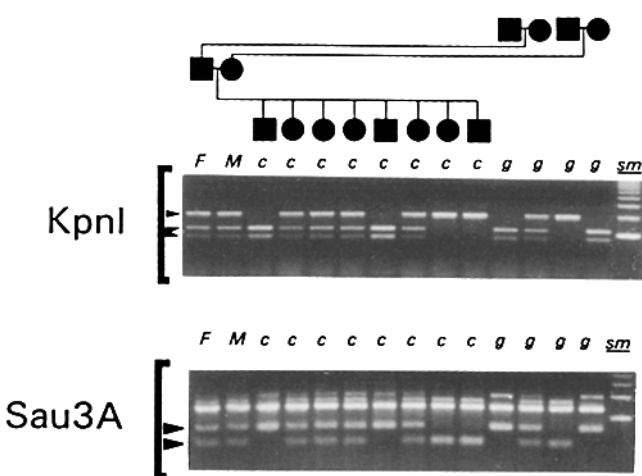


Figure 2. Mendelian segregation of the KpnI and Sau3A V β 18 DNA polymorphisms. The V β 18 gene was amplified using primers V β 18-3 and -4 from the individuals in a three-generation pedigree (F, father; M, mother; c, child; g, grandparent). The PCR products were digested with either the KpnI or Sau3A restriction enzyme and genotypes were determined based on the polymorphic DNA fragments observed. The pointers on the left indicate the allelic restriction fragments, with the joined pointers (after KpnI digestion) indicating a single allele (though the lower fragment is noticeably faint in the heterozygotes). Sau3A digestion also resulted in a monomorphic DNA fragment present in all individuals. On the far right, sm indicates the 123-bp ladder used as a size marker.

V β 18 polymorphisms were determined for a larger group of individuals. First, according to the DNA sequences surrounding the polymorphisms at nucleotide positions 524 and 620 (Fig. 1), these polymorphisms would result in altering restriction enzyme recognition sites. The base change at position 524 alters a KpnI restriction site, and the DNA polymorphism at position 620 changes a Sau3A restriction site, as demonstrated in Fig. 2. Genotypes for all three V β 18 polymorphisms were obtained using the semi-automated OLA

typing system. Analysis of the inheritance patterns for these three DNA polymorphisms revealed proper Mendelian segregation, as shown in Fig. 3. The segregation of the V β 18 genotypes was completely compatible with cosegregation (i.e., no recombinants) with haplotypes of the TCR β chain previously determined in these individuals (6).

Within our tested population (152 chromosomes), the nucleotide changes at positions 524 and 620 had identical allele frequencies of 68% for the C substitution and 32% for the T substitution. 11% of the individuals tested were homozygous for the T substitution at nucleotide 524, and therefore, based on the presence of the stop codon, these 11% of the individuals would not produce a functional V β 18 protein product. The allele frequency for the silent base substitution at nucleotide 661 was slightly different from those found for the other two polymorphisms (524 and 620), i.e., at position 661 the allele frequency for the C substitution was 53% and the T substitution was 47%. Even though one of the substitutions, 524, leads to a stop codon, the genotype distributions for all three polymorphisms were consistent with Hardy-Weinberg equilibrium.

Linkage Disequilibrium Analysis. We statistically examined the extent of association between alleles of the V β 18 polymorphism(s) and alleles of other previously reported TCR β DNA polymorphisms (Table 2). Because of complete linkage disequilibrium among the three point mutations within the V β 18 gene ($D' = 1.00$), only three DNA haplotypes were observed (as shown in Fig. 1). The extent of linkage disequilibrium was also measured between the V β 18 polymorphisms and other biallelic polymorphisms previously studied in this same population of CEPH parents (6, 14). Although the alleles of the V β 18 polymorphisms did not show statistical association with other V β gene polymorphisms (Table 2), polymorphisms at the 3' end of the TCR β gene complex (C β 2 region) were significantly nonrandomly associated with the 524 and 620 V β 18 polymorphism(s) ($p < 0.0001$).

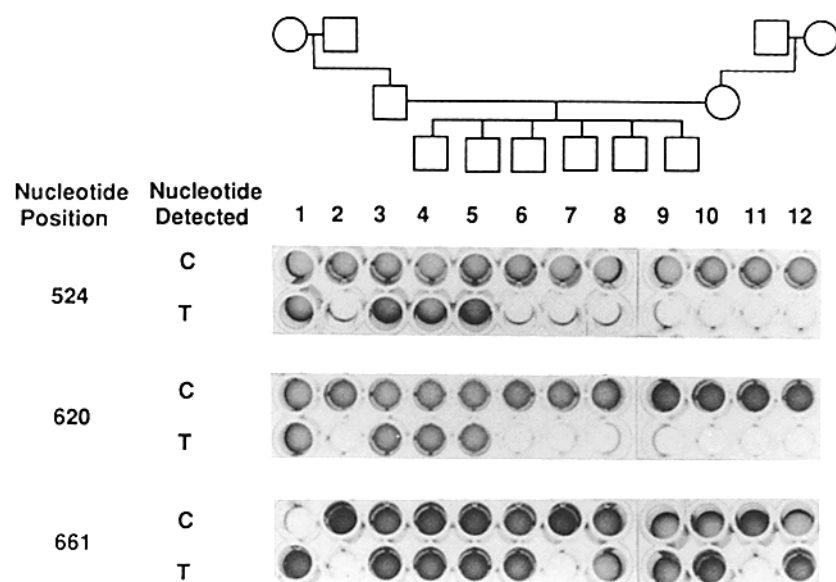


Figure 3. A composite figure assembled from photographs of microtiter plates containing OLA genotypes that show Mendelian segregation of the three V β 18 sequence polymorphisms. Microtiter wells containing the reporter group digoxigenin form a colored product (dark wells) and indicate a perfect nucleotide match between the ligating probes and the amplified DNA target.

Table 1. DNA Sequences for the Amplification Primers and Ligation Probes Used in Typing the V β 18 DNA Polymorphisms

Amplification primers	Sequences	
Vβ18-1	GAGTGCACCTGTGGAGGGAA	
Vβ18-2	TCACATTGGCCAGCAGTCAG	
Vβ18-3	ATTCATCAATGGCCAGCGAC	
Vβ18-4	GGAGCTTCTTAGAACTCAG	
<u>Ligation probes*</u>		
Vβ18-524	B-CCCCAACCTATACTGGTACC B-CCCCAACCTATACTGGTACT	pGACAGGCTGCAGGCAGGG-D
Vβ18-620	B-GAATCTCTCAGCCTCCAGAC B-GAATCTCTCAGCCTCCAGAT	pCCCAGGACCGGCAGTTCAT-D
Vβ18-661	B-TGAGTTCTAAGAAGCTCCTC B-TGAGTTCTAAGAAGCTCCTT	pCTCAGTGACTCTGGCTTCTAT-D

* B, biotin; p, phosphorylated; D, digoxigenin.

Table 2. Analysis of Linkage Disequilibrium between V β 18 Polymorphisms and Other Polymorphisms in the TCR β Gene Complex

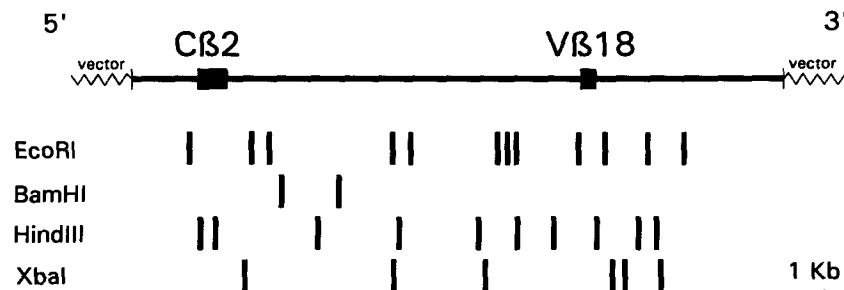
V β 18-524/620* and:	Haplotypes	D' [†]	Q (χ^2) [§]	V β 18-661 and:	Haplotypes	D'	Q (χ^2)
V β 1	140	0.04	0.1	V β 1	130	0.07	0.1
V β 8	116	0.18	1.3	V β 8	92	-0.20	2.9
V β 11	114	-0.23	1.9	V β 11	94	-0.20	2.5
C β 2-1	118	-0.28	2.2	C β 2-1	102	-0.30	4.5
C β 2-5	102	-1.0	31.4	C β 2-5	94	-0.35	8.7
C β 2-6	108	0.69	24.8	C β 2-6	98	0.27	6.4
C β 2-7	98	-1.0	27.2	C β 2-7	92	-0.37	9.0
V β 18-661	84	1.0	31.2	V β 18-524/620*	84	1.0	31.2

The order of the polymorphisms compared in this table (top to bottom) is the same as the physical order of the polymorphisms along the chromosome (centromere to telomere) (6, 14).

* The V β 18 position 524 and 620 polymorphisms are synonymous for this analysis since there was a singular correspondence between alleles at these loci (haplotypes, 86; D', 1.0; Q, 86.0).

[†] For D', a value of 1.00 (or -1.00) indicates maximum linkage disequilibrium; a value of 0.0 would indicate complete random association between the loci.

[§] For the Q statistic (χ^2 distribution, 1-degree freedom), Q >15 corresponds to $p < 0.0001$, which rejects the null hypothesis of random association between the alleles at that pair of loci.

**Figure 4.** Restriction map of the cosmid insert containing V β 18 and C β 2 genes. Cosmid H29 was shown by hybridization to contain both the V β 18 and C β gene segments. A restriction map of the cosmid was constructed by partial digestion and hybridization with either the gene segments or vector sequences. Restriction sites for EcoRI, BamHI, HindIII, and XbaI are shown as vertical lines below the cosmid.

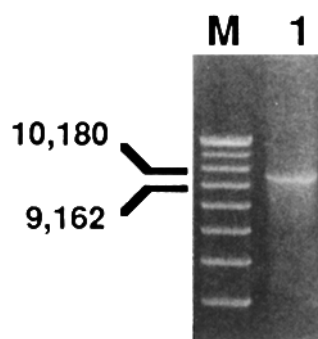


Figure 5. Ethidium bromide-stained gel containing the amplified product from the H29 cosmid using a primer located at the 3' end of the C β 2 gene and one located 3' of the V β 18 gene. Lane M contains a size marker consisting of a 1-kb ladder. The sizes of the two size marker bands flanking the amplified C β 2-V β 18 product are indicated to the left.

Physical Mapping of the V β 18 Cosmid. Based on the linkage disequilibrium results (Table 2), we sought to confirm the suggested 3' location of the V β 18 gene relative to C β 2. The H29 cosmid that contained V β 18 was tested for hybridization to 22 other V β families and C β . Only hybridization to a C β probe was detected. PCR was used to confirm that this cosmid contained both the V β 18 and C β genes. Physical mapping of the H29 cosmid with four restriction enzymes and hybridization with V β 18 and C β probes indicated that the V β 18 gene was located within 15 kb of the C β 2 gene (Fig. 4). Previous studies have shown that the mouse V β 14 gene (gene counterpart of human V β 18) has an opposite transcriptional orientation to that of other V β genes as well as the D, J, and C gene segments (20). By using PCR with the H29 cosmid, we have determined the transcriptional orientation of human V β 18 relative to the C β 2 gene segment. DNA amplification with a primer located just 3' of the end of the C β 2 gene, in combination with a primer located just downstream of the V β 18 gene, indicated that the V β 18 gene segment is located \sim 9 kb 3' of the C β 2 gene (Fig. 5). Since these PCR primers must face each other to amplify the intervening DNA sequence, these data also indicate that the transcriptional orientation for human V β 18 is opposite to that of the C β 2 gene, and therefore similar to the orientation of the mouse V β 14 and C β 2.

It is worth noting that several different pairs of amplification primers from the 3' end of the C β 2 gene and the 3' end of the V β 18 gene could be used to obtain a similarly sized PCR product from cosmid DNA (data not shown). In addition, we were also able to sequence the ends of the 9.5-kb PCR product (Fig. 5) in low-melt agarose (see Materials and Methods), which verified the origins of this PCR product, i.e. C β 2 at the 5' end and V β 18 at the 3' end (data not shown).

V β 18 RNA Expression in Peripheral Blood. As a result of the null allele for V β 18 in the population, we sought to confirm the importance of this polymorphism in the expression level of V β 18 in individuals homozygous for the V β 18 stop codon. Several members from three families were tested by KpnI restriction enzyme analysis of genomic DNA for the V β 18 null allele (KpnI⁻). Representatives of each of the three genotypes were identified (Fig. 6 A) and PCR was used to amplify the V β 18 RNA (cDNA) from all these individuals. Based on the intensity of the V β 18 PCR product, and the control C α product in each lane, V β 18-C β message was

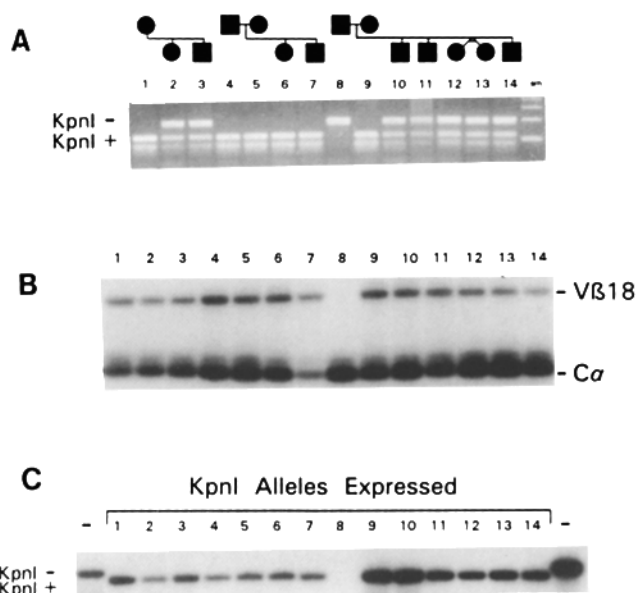


Figure 6. V β 18 DNA genotypes and RNA expression in PBL of 14 individuals. (A) PCR-amplified V β 18 gene segment from genomic DNA was digested with KpnI. The DNA polymorphism that results in the stop codon also destroys this KpnI site. The family structures are shown above. (B) The somatically rearranged V β 18 gene was amplified from the PBL RNA (cDNA) of these same individuals. The forward primer V β 18-3 and a reverse C β primer (5'-AGTCATTGAGGGCGGGCTG-3') produced the larger V β 18-containing product, and each tube also contained a TCR C α primer pair (5'-CAGCATTTATCCAGAAGACAC-3', and 5'-CCTCAGCTGGACCACAGC-3') as a positive control for amplification. (C) The rearranged V β 18 gene was amplified from cDNA by PCR using the forward primer V β 18-3 and a reverse C β primer (5'-GGCAGACAGGACCCCTTG-3'); the C α primer pair listed above was also included. The PCR products were digested with KpnI restriction enzyme to confirm that the V β 18 null allele (no KpnI site) was not expressed. As a control for the V β 18-C β KpnI⁻ allele, the outer lanes (-) contain the PCR product to which KpnI enzyme was not added. The C α PCR-positive controls in each lane are not shown.

detected in the peripheral blood of all individuals, except the person homozygous for the stop codon (Fig. 6 B). As even stronger evidence of the relationship between the stop codon and V β 18 expression, the amplified V β 18-C β message was digested with KpnI (Fig. 6 C). The results show that in the eight individuals studied who possessed the null allele (either heterozygous or homozygous individuals), only the V β 18 KpnI⁺ allele was expressed at detectable levels in the peripheral blood.

Discussion

Allelic variation in the germline DNA of the TCR genes has been suspected to influence the expressed TCR repertoire by altering amino acids that change the TCR's ability to recognize antigen and/or MHC. DNA polymorphisms in TCR genes could change the expressed T cell repertoire either qualitatively or quantitatively through the process of clonal selection and expansion in the thymus or periphery. It is also possible that DNA polymorphisms could influence the quan-

tity of an expressed TCR protein by altering regulatory elements, or through translational changes such as creating a stop codon. The potential for functional relevance of TCR polymorphisms has been the basis for many disease association studies that have compared the frequency of various TCR DNA polymorphisms in autoimmune and control populations (reviewed in references 1 and 9).

In this study, we describe an extreme example of a functionally relevant DNA polymorphism that results in the presence of a stop codon in the V β 18 gene segment. This polymorphism was present in 54% of the population that we studied. Importantly, 11% of the individuals tested were homozygous for this null allele, indicating a lack of cell surface expression of the V β 18 TCR protein. Coincidentally, there were two other DNA polymorphisms in the V β 18 gene, one of which would have resulted in an amino acid change from proline to serine, in the absence of the stop codon.

The observation of complete allelic association between the V β 18 polymorphisms could have implications in terms of the origins of these polymorphisms. Since the V β 18 alleles at positions 524 and 620 were correlated completely, this could be due to the simultaneous "creation" of the polymorphisms in a single ancestral genetic event. A more speculative but immunological explanation for the complete allelic correlation is evolutionary selection at the protein level since the alleles of both polymorphisms would affect expression and possibly structure. At present time, whenever the serine residue is coded for at the second polymorphic site, the termination codon is encoded at the first polymorphic site. The serine/proline substitution site would occur within the fourth hypervariable region in the TCR β chain (21) and might be involved in antigen/MHC contact. Based on the location of this amino acid substitution, and the radical nature of the substitution (22), generous speculation might suggest that the V β 18 haplotype with the stop codon and serine residue has been selected to prevent protein expression of the V β 18 with a serine residue at this position.

The strong disequilibrium between the C β -KpnI site (C β 2-5; Table 2) and the V β 18 polymorphisms has relevance to disease association studies since previous studies that have used the C β -KpnI RFLP have therefore indirectly measured the V β 18 polymorphism association with systemic lupus erythematosus, multiple sclerosis, and myasthenia gravis (23, 24). Also, the lack of any detectable disequilibrium between the V β 18 polymorphisms and the C β -BglII RFLP (C β 2-1), as well as the other more distant 5' V β RFLPs (Table 2), implies that the many positive disease associations found with those RFLPs (1, 9) cannot be directly explained by the V β 18 stop codon. Vice versa, studies reporting no disease association with the previously reported V β and C β -BglII RFLPs in Table 2 (except C β -KpnI) cannot conclude that the V β 18 polymorphisms are not involved in the disease susceptibility.

Physical mapping of the human and mouse TCR β gene complexes has shown a remarkable conservation in the organization of homologous V β genes between these two species (19, 25). The human V β 18 gene is most similar to the mouse V β 14 gene based on the high nucleic acid homology between these gene segments (77%). Therefore, since the

mouse V β 14 has been shown to be the only V gene located 3' of the TCR β C region (26), it was predicted that the human V β 18 gene would be located 3' of the C region (19). This was suggested by linkage disequilibrium (Table 2) and confirmed by isolating a cosmid that contained both the human V β 18 and C region C β 2 gene. Restriction enzyme and PCR mapping of this cosmid showed V β 18 to be \sim 9 kb 3' of the C β 2 gene segment. This 9-kb distance in humans is similar to the 10-kb distance between C β 2 and V β 14 in mice (26). Multiple PCR analyses also showed that the human V β 18 has an opposite transcriptional polarity relative to the TCR β D, J, and C region genes and other V β genes, as shown for mouse V β 14 (26). Thus, it may similarly be expected that human V β 18 undergoes a chromosomal inversion to juxtapose V β 18 with the D, J, and C region genes before successful transcription.

Since the human V β 18 gene was originally cloned from an acute lymphocytic leukemia cell line (20), and since it is 3' of C β as well as being polymorphic for a stop codon, it may be questioned as to the quantitative contribution of a functional V β 18 protein product to the total TCR β repertoire. Choi et al. (27) estimated that V β 18 (V β 20 in their nomenclature) accounted for \sim 3.4% of the total V β expression after anti-CD3 stimulation of peripheral T cells from one individual, while another group failed to detect V β 18 expression in peripheral blood from one donor (28). V β 18 is a single member subfamily (29), so there are no other V β 18 subfamily members that might subsume the role of V β 18 and prevent this hole in the TCR repertoire caused by the null allele.

Our result showing greatly reduced (undetectable) V β 18 RNA expression in the peripheral blood of the null allele in all eight individuals who possessed the null allele (Fig. 6, *c* and *b*) is consistent with the mechanisms of T cell selection that occur in the thymus. The positive or negative selection of T cells in the thymus is dependent on the specific α and β proteins on the cell's surface. T cells that rearrange and transcribe the nonfunctional V β 18 gene segment could not be positively selected because they would be unable to assemble a TCR α/β protein heterodimer. Thus, these V β 18-rearranged cells could not be expected to populate the peripheral blood and immune system to an appreciable level. In an individual homozygous for the V β 18 null allele, a very low level of rearranged V β 18 message in PBL could be accounted for by the infrequent additional rearrangement that can occur on the other chromosome 7 homologue in T cells that express some other V β gene on their cell surface. The failure to detect V β 18-C β RNA message from the null allele is also consistent with what has been previously seen in mice with a similar V β stop codon (30, and see below [BALB/c V β 19]).

While other polymorphic stop codons have not yet been reported in human TCR genes, there are two examples of polymorphic stop codons in mouse TCR V β genes. Mouse V β 19 is expressed in the SJL strain, but the BALB/c strain has deleted a single base in the leader exon that results in a frameshift mutation (30). A second example of a polymorphic stop codon in a mouse TCR β gene is mouse V β 17. Certain

strains of mice possess a TCR β haplotype that encodes a V β 17 gene with a single mutation in the coding region that results in a stop codon (31). Other strains of mice with a functional V β 17 gene have their V β 17-bearing T cells clonally eliminated in those mice that express a certain self-superantigen and the MHC class II molecule IE (32). Wade et al. (31) speculated that although self-tolerance mechanisms would seem to be able to control such autoimmune cells reactive to the self-superantigen, perhaps evolutionary pressures have acted

to provide an additional means of eliminating such autoreactive T cells from a portion of the population, via a stop codon. It is therefore interesting to note that the T cells bearing human V β 18 appear to be stimulated by the *Staphylococcus aureus* enterotoxin superantigens (27). This parallel between these mouse and human V β genes allows similar speculation on whether past selective pressures on humans could have had an importance in the prevalence of the polymorphic stop codon in the human V β 18 gene.

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Note added in proof: Malhotra et al. (33) have recently reported the detection, by a different approach and technology, of the mutation at nucleotide position 524 in the V β 18 gene (the V β 20 gene according to their nomenclature).

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